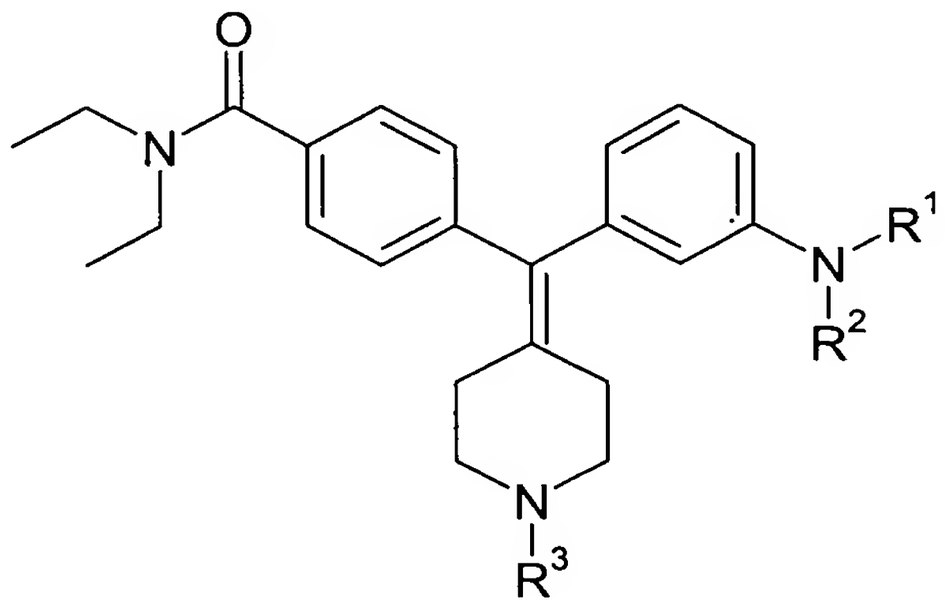
Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Original) A compound of formula I, or a pharmaceutically acceptable salt thereof:



I

wherein

R^1 is selected from C_{3-6}alkyl, C_{6-10}aryl, C_{2-9}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-9}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, C_{3-10}cycloalkyl-C_{1-4}alkyl, R^8-C(=O)-, R^8-$S(=O)_2$-, R^8-S(=O)-, R^8-NHC(=O)-, R^8-C(=S)- and R^8-NH-C(=S)-, wherein R^8 is selected from C_{3-6}alkyl, C_{6-10}aryl, C_{2-9}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-9}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, and C_{3-10}cycloalkyl-C_{1-4}alkyl, wherein said C_{3-6}alkyl, C_{6-10}aryl, C_{2-9}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-9}heteroaryl-C_{1-4}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl used in defining R^1 and R^8 are optionally substituted with one or more groups selected from -R, $–NO_2$, -OR, -Cl, -Br, -I, -F, $-CF_3$, -C(=O)R, -C(=O)OH, $-NH_2$, -SH, -NHR, $-NR_2$, -SR, $-SO_3H$, $-SO_2R$, -S(=O)R, -CN, -OH, -C(=O)OR, $-C(=O)NR_2$, -NRC(=O)R, and -NRC(=O)-OR, wherein R is, independently, selected from -H, C_{1-6}alkyl and phenyl;

R^2 is selected from -H and C_{1-6}alkyl optionally substituted with one or more groups selected from halogen, $-CF_3$, -OH, C_{1-3}alkoxy, and halogen, or R^1 and R^2 are C1-3alkylene that together form a portion of a ring; and

R^3 is selected from -H, C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl, wherein said C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl,

and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-6}alkyl, halogenated C_{1-6}alkyl, -NO_2, -CF_3, C_{1-6}alkoxy and halogen.

2. (Original) A compound according to claim 1, wherein

R^1 is selected from C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, C_{3-10}cycloalkyl-C_{1-4}alkyl, wherein said C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, C_{3-10}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-4}alkyl, halogen, -CF_3, -OH, C_{1-3}alkoxy, phenoxy, and halogen;

R^2 is selected from –H and C_{1-3}alkyl; and

R^3 is selected from –H and C_{1-6}alkyl-O-C(=O)-.

3. (Original) A compound according to claim 2,

wherein R^1 is R^9-CH_2-, wherein R^9 is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, triazolyl, pyrrolyl, thiazolyl, N-oxido-pyridyl, benzyl, pyridylmethyl, thienylmethyl, furylmethyl, imidazolylmethyl, triazolylmethyl, pyrrolylmethyl, thiazolylmethyl and N-oxido-pyridylmethyl, optionally substituted with one or more groups selected from C_{1-4}alkyl, halogen, -CF_3, -OH, C_{1-3}alkoxy, phenoxy and halogen; and

R^2 and R^3 are hydrogen.

4. (Original) A compound according to claim 3,

wherein R^9 is selected from benzyl, phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl and thiazolyl, optionally substituted with one or more groups selected from C_{1-4}alkyl, halogen, -CF_3, -OH, C_{1-3}alkoxy, phenoxy, and halogen.

5. (Original) A compound according to claim 4, wherein

wherein R^9 is selected from benzyl, phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl and thiazolyl.

6. (Original) A compound according to claim 1, wherein

R^1 is selected from C_{3-6}alkyl, C_{3-10}cycloalkyl, and C_{3-10}cycloalkyl-C_{1-4}alkyl, wherein said C_{3-6}alkyl, C_{3-10}cycloalkyl, and C_{3-10}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-4}alkyl, halogen, -CF_3, -OH, C_{1-3}alkoxy, phenoxy, and halogen;

R^2 is –H or C_{1-3}alkyl; and

R^3 is –H, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl, wherein said C_{1-6}alkyl, C_{3-6}cycloalkyl, C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-4}alkyl, halogen, -CF_3, -OH, C_{1-3}alkoxy, phenoxy, and halogen.

7. (Original) A compound according to claim 6, wherein

R^1 is selected from 1-propyl, 2-propyl, 1-butyl, 2-butyl, t-butyl, 2-methyl-1-propyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, and cyclononyl;

R^2 is selected from –H, methyl, ethyl, 1-propyl and 2-propyl; and

R^3 is selected from –H, methyl, ethyl, allyl, 3,3-dimethyl-allyl, cyclopropylmethyl, 2-methoxy-ethyl, and 3-methoxy-1-propyl.

8. (Original) A compound according to claim 1, wherein

R^1 is selected from R^8-C(=O)-, R^8-$S(=O)_2$-, R^8-S(=O)-, R^8-NHC(=O)-, R^8-C(=S)- and R^8-NH-C(=S)-, wherein R^8 is selected from C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, and C_{3-10}cycloalkyl-C_{1-4}alkyl; wherein said C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with C_{1-4}alkyl, halogen, -CF_3, -OH, C_{1-3}alkoxy, phenoxy, and halogen;

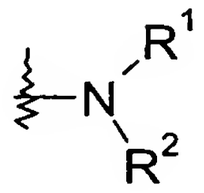
R^2 is –H; and

R^3 is selected from –H and C_{1-6}alkyl-O-C(=O)-.

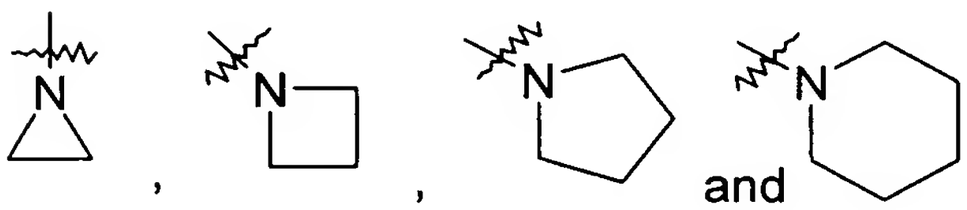
9. (Original) A compound according to claim 8, wherein

R^8 is selected from phenyl, benzyl, phenethyl and cyclohexyl, wherein said phenyl, benzyl, phenethyl and cyclohexyl are optionally substituted with one or more groups selected from methyl, methoxy and halogen.

10. (Original) A compound according to claim 1, wherein



of formula I is selected from



; and

R^3 is selected from –H and $C_{1\text{-}6}$alkyl-O-C(=O)-.

11. (Original) A compound selected from:

1) 4-[[3-(benzylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

2) N,N-diethyl-4-[{3-[(3-furylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,

3) N,N-diethyl-4-(piperidin-4-ylidene{3-[(thien-3-ylmethyl)amino]phenyl}methyl)benzamide,

4) N,N-diethyl-4-[{3-[(2-phenylethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,

5) 4-[{3-[(4-chlorobenzyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

6) N,N-diethyl-4-[piperidin-4-ylidene(3-{[3-(trifluoromethyl)benzyl]amino}phenyl)methyl]benzamide,

7) 4-[{3-[(2-chlorobenzyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

8) N,N-diethyl-4-[piperidin-4-ylidene(3-{[4-(trifluoromethyl)benzyl]amino}phenyl)methyl]benzamide,

9) N,N-diethyl-4-[{3-[(2-furylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,

10) N,N-diethyl-4-(piperidin-4-ylidene{3-[(thien-2-ylmethyl)amino]phenyl}methyl)benzamide,

11) 4-[{3-[(cyclohexylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

12) N,N-diethyl-4-{piperidin-4-ylidene[3-(propylamino)phenyl]methyl}benzamide,

13) 4-[[3-(cyclohexylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

14) 4-[[3-(cyclopentylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

15) 4-[[3-(cycloheptylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

16) 4-[{3-[cyclopentyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

17) 4-[[3-(benzoylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

18) N,N-diethyl-4-[{3-[(phenylacetyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,

19) 4-[{3-[(cyclohexylcarbonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

20) 4-[{3-[(cyclohexylacetyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

21) 4-[(3-{[(2-chlorophenyl)acetyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

22) 4-[(3-{[(3-chlorophenyl)acetyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

23) N,N-diethyl-4-[(3-{[(5-methylthien-2-yl)acetyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,

24) 4-[(3-{[(5-chlorothien-2-yl)acetyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

25) N,N-diethyl-4-[(3-{[(2S)-2-phenylpropanoyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,

26) N,N-diethyl-4-[(3-{[(2R)-2-phenylpropanoyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,

27) N,N-diethyl-4-[(3-{[(2S)-2-phenylbutanoyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,

28) N,N-diethyl-4-[(3-{[(2R)-2-phenylbutanoyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,

29) 4-[{3-[benzoyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

30) 4-[{3-[(anilinocarbonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

31) 4-[(3-{[(benzylamino)carbonyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

32) N-{3-[{4-[(diethylamino)carbonyl]phenyl}(piperidin-4-ylidene)methyl]phenyl}piperidine-1-carboxamide,

33) N,N-diethyl-4-[{3-[(phenylsulfonyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,

34) 4-[{3-[(benzylsulfonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

35) 4-[(3-anilinophenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

36) N,N-diethyl-4-[{3-[methyl(phenyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,

37) N,N-diethyl-4-[{3-[ethyl(phenyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,

38) N,N-diethyl-4-[(3-{[(1S)-1-phenylethyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,

39) N,N-diethyl-4-[(3-{[(1R)-1-phenylethyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,

40) 4-[(3-{[(1R)-1-cyclohexylethyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

41) 4-[(3-{[(1S)-1-cyclohexylethyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

42) N,N-diethyl-4-[{3-[(1-methyl-1-phenylethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,

43) 4-[{3-[cyclohexyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

44) N,N-diethyl-4-[piperidin-4-ylidene(3-piperidin-1-ylphenyl)methyl]benzamide,

45) N,N-diethyl-4-[piperidin-4-ylidene(3-pyrrolidin-1-ylphenyl)methyl]benzamide,

46) N,N-diethyl-4-[[3-[(2-ethyl-1-oxobutyl)amino]phenyl]-4-piperidinylidenemethyl]-benzamide,

47) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-1-methyl-1H-1,2,3-benzotriazole-5-carboxamide,

48) 6-chloro-N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-3-pyridinecarboxamide,

49) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-methoxy-benzamide,

50) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-quinoxalinecarboxamide,

51) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2,5-difluoro-benzamide,

52) 3-chloro-N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-thiophenecarboxamide,

53) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-3-methyl-benzamide,

54) N,N-diethyl-4-[[3-[[(methylphenylamino)carbonyl]amino]phenyl]-4-piperidinylidenemethyl]-benzamide, and pharmaceutically acceptable salts thereof.

12-13. (Cancelled)

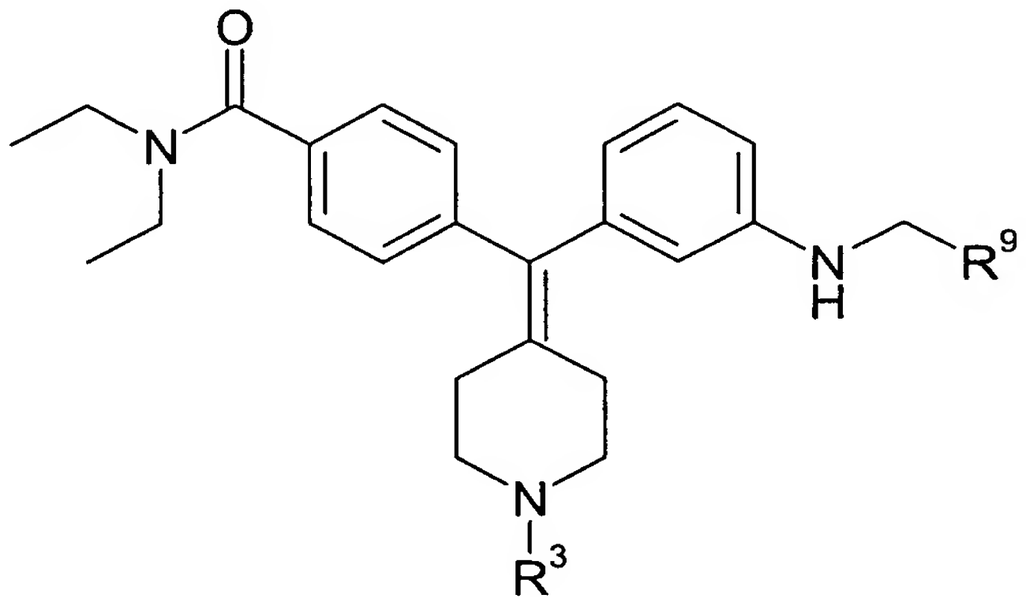
14. (Currently Amended) A pharmaceutical composition comprising a compound according to ~~any one of claims 1-11~~claim 1 and a pharmaceutically acceptable carrier.

15. (Currently Amended) A method for the therapy of pain in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to ~~any one of claims 1-11~~claim 1.

16. (Currently Amended) A method for the therapy of functional gastrointestinal disorders in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to ~~any one of claims 1-11~~claim 1.

17. (Currently Amended) A method for the therapy of anxiety in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to ~~any one of claims 1-11~~claim 1.

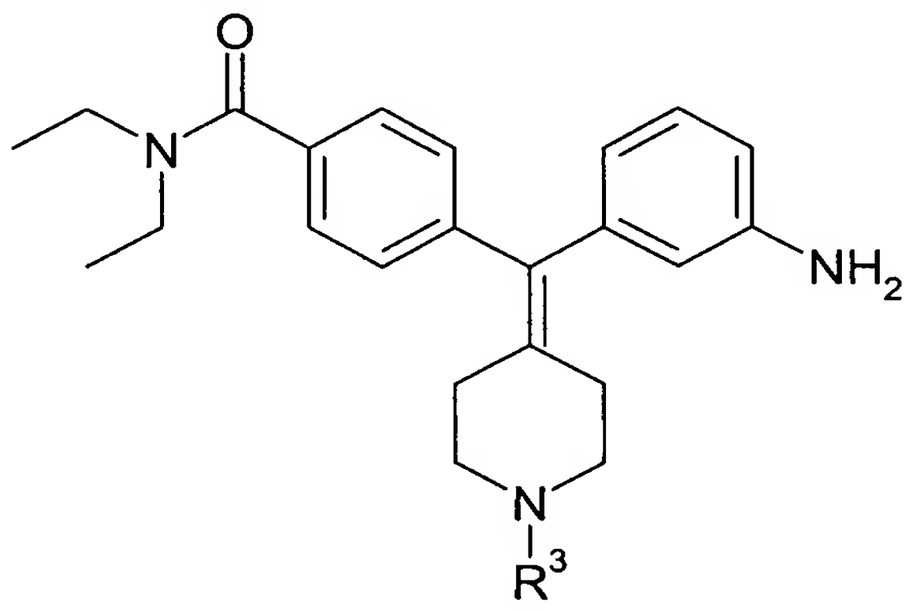
18. (Original) A process for preparing a compound of formula III,



III

comprising:

reacting a compound of formula II,



II

with R^9-CHO in the presence of a reducing agent to form the compound of formula III,

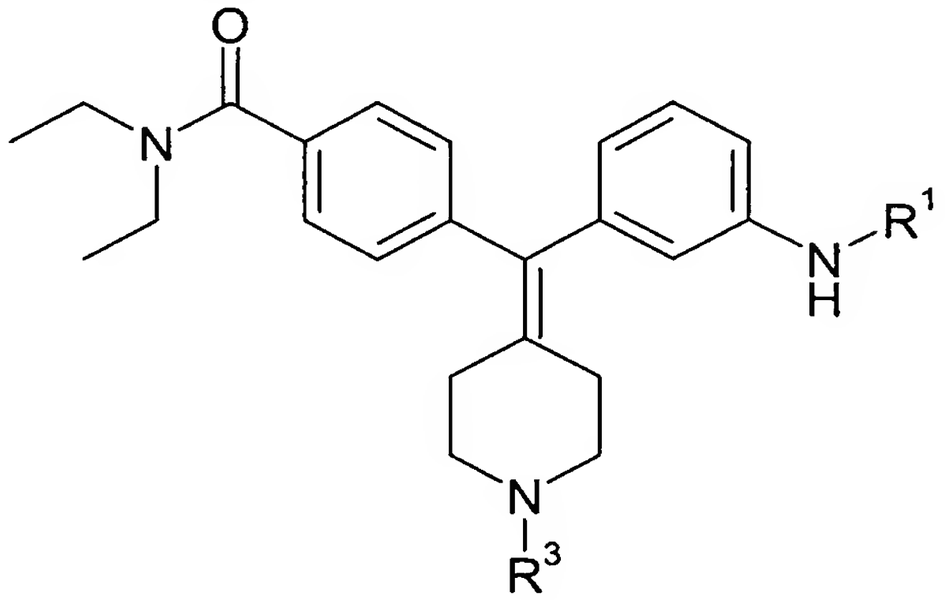
wherein

R^9 is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, triazolyl, pyrrolyl, thiazolyl, N-oxido-pyridyl, benzyl, pyridylmethyl, thienylmethyl, furylmethyl, imidazolylmethyl, triazolylmethyl, pyrrolylmethyl, thiazolylmethyl and

N-oxido-pyridylmethyl, optionally substituted with one or more groups selected from C_{1-4}alkyl, halogen, $-CF_3$, -OH, C_{1-3}alkoxy, phenoxy and halogen; and

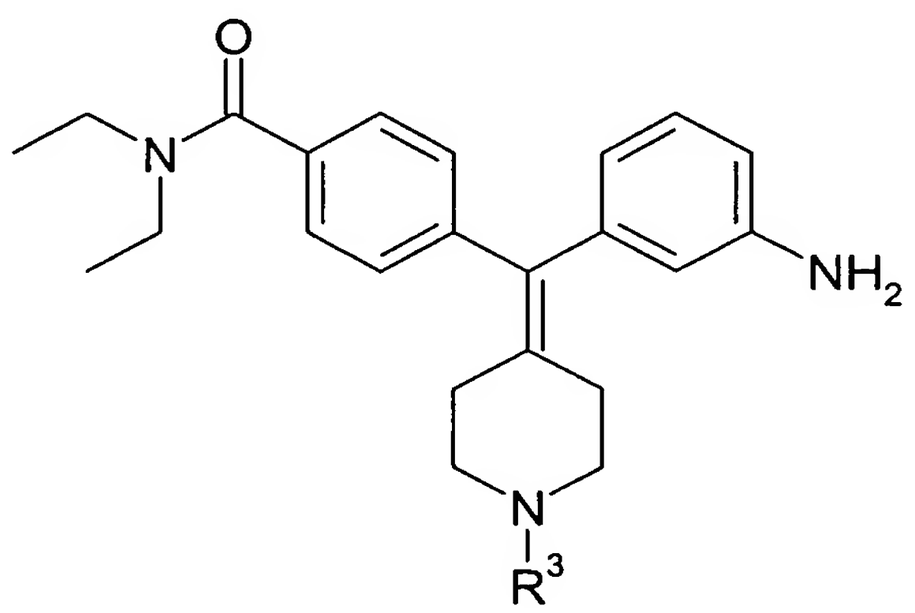
R^3 is selected from C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl, wherein said C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-6}alkyl, halogenated C_{1-6}alkyl, $-NO_2$, $-CF_3$, C_{1-6}alkoxy and halogen.

19. (Original) A process for preparing a compound of formula IV,



IV

comprising: reacting a compound of formula II,



II

with R^1-X to form the compound of formula IV,

wherein

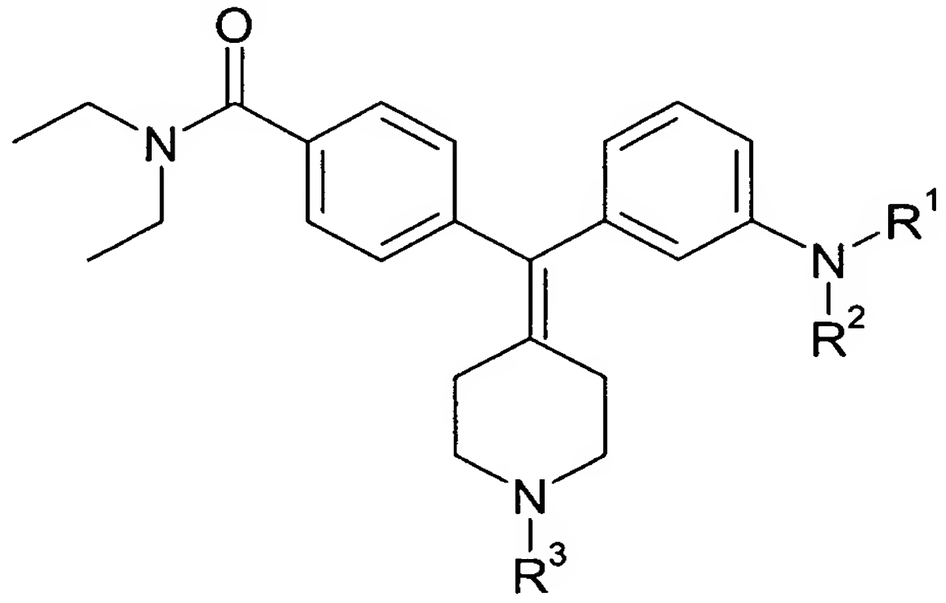
X is halogen;

R^1 is selected from C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, C_{3-10}cycloalkyl-C_{1-4}alkyl, wherein said C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl,

C_{3-10}cycloalkyl, C_{3-10}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-4}alkyl, halogen, $-CF_3$, -OH, C_{1-3}alkoxy, phenoxy, and halogen; and

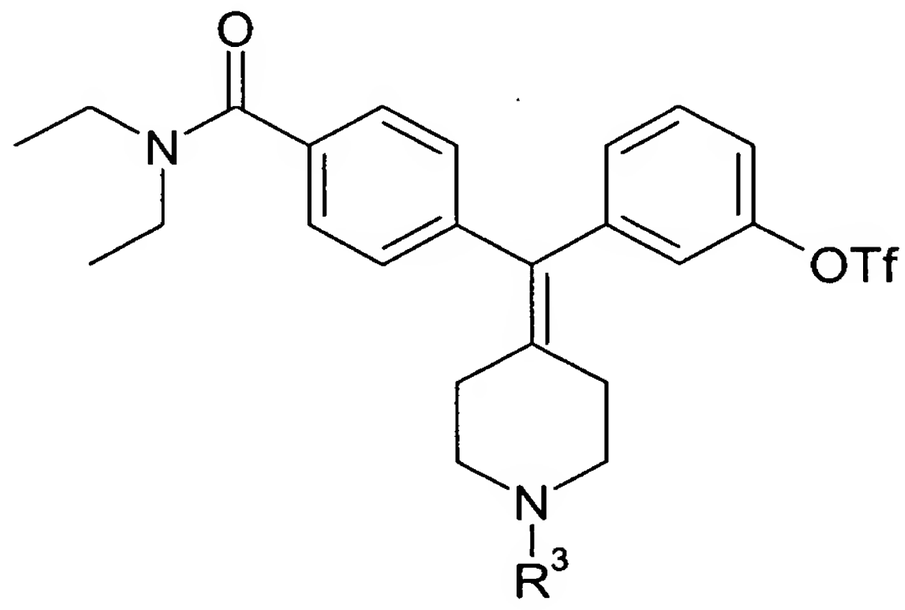
R^3 is selected from C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl, wherein said C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-6}alkyl, halogenated C_{1-6}alkyl, $-NO_2$, $-CF_3$, C_{1-6}alkoxy and halogen.

20. (Original) A process for preparing a compound of formula I,



I

comprising: reacting a compound of formula V,



V

with R^1R^2NH to form the compound of formula I,

wherein

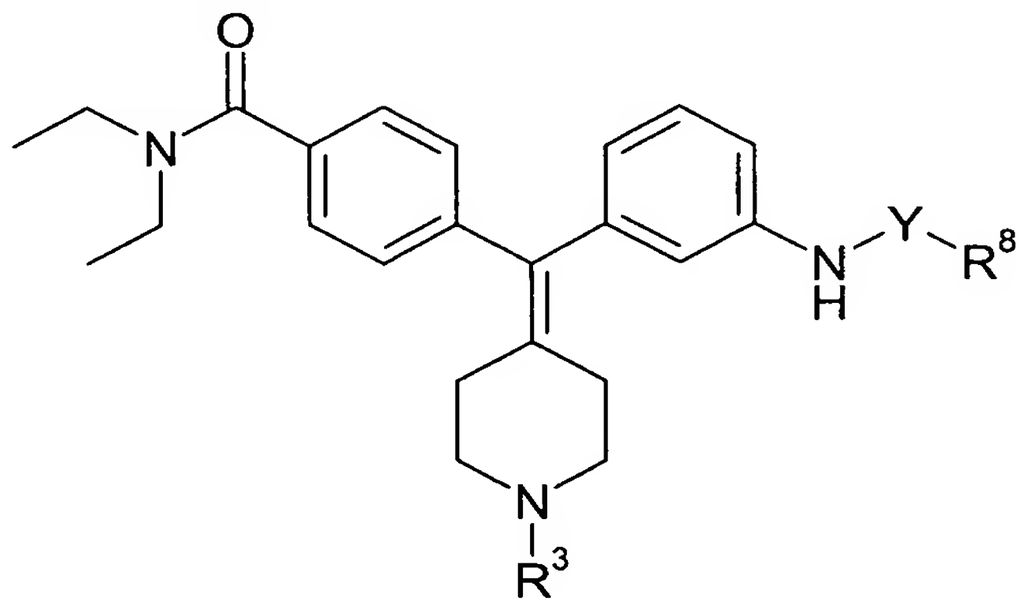
R^1 is selected from C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, C_{3-10}cycloalkyl-C_{1-4}alkyl, wherein said C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl,

C_{3-10}cycloalkyl, C_{3-10}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-4}alkyl, halogen, $-CF_3$, -OH, C_{1-3}alkoxy, phenoxy, and halogen;

R^2 is selected from -H and C_{1-6}alkyl optionally substituted with one or more groups selected from halogen, $-CF_3$, -OH, C_{1-3}alkoxy, and halogen, or R^1 and R^2 are C1-3alkylene that together form a portion of a ring; and

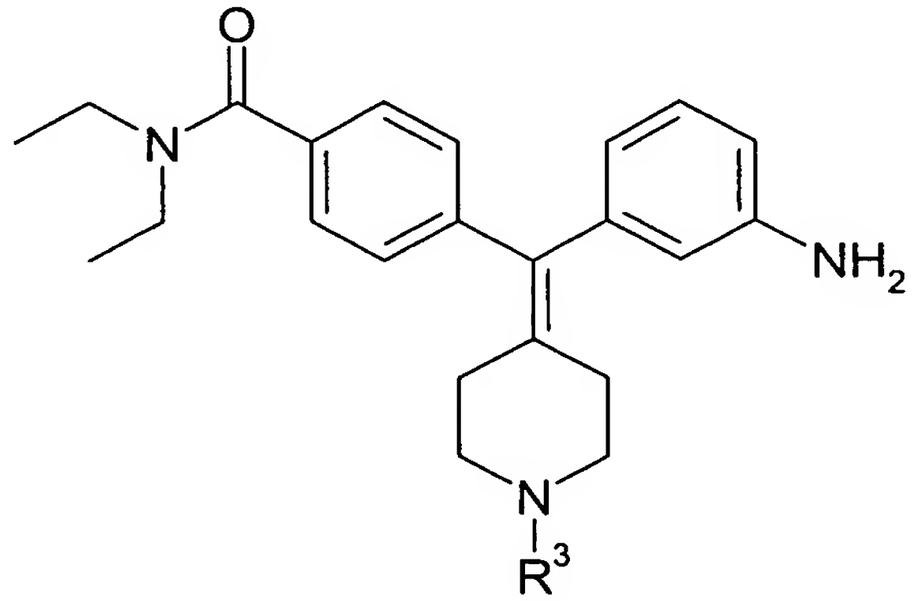
R^3 is selected from C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl, wherein said C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-6}alkyl, halogenated C_{1-6}alkyl, $-NO_2$, $-CF_3$, C_{1-6}alkoxy and halogen.

21. (Original) A process for preparing a compound of formula VI,



VI

comprising: reacting a compound of formula VII,



VII

with R^8-Y-X or R^8-Y-O-Y-R^8 to form the compound of formula VI:

wherein

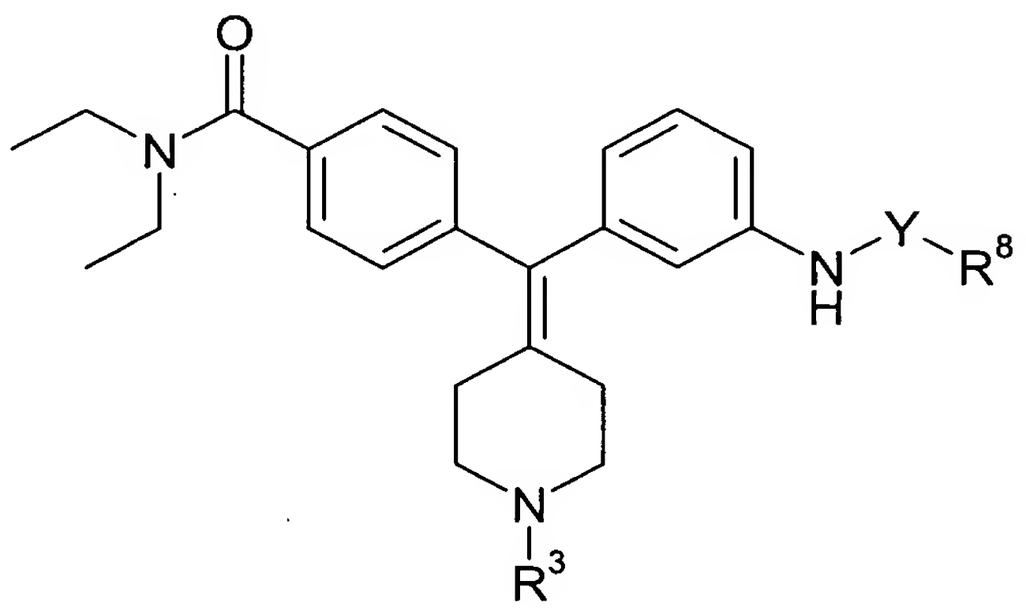
X is halogen;

Y is selected from –C(=O)- and –$S(=O)_2$-;

R^8 is selected from C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, and C_{3-10}cycloalkyl-C_{1-4}alkyl; wherein said C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with C_{1-4}alkyl, halogen, -CF_3, -OH, C_{1-3}alkoxy, phenoxy, and halogen; and

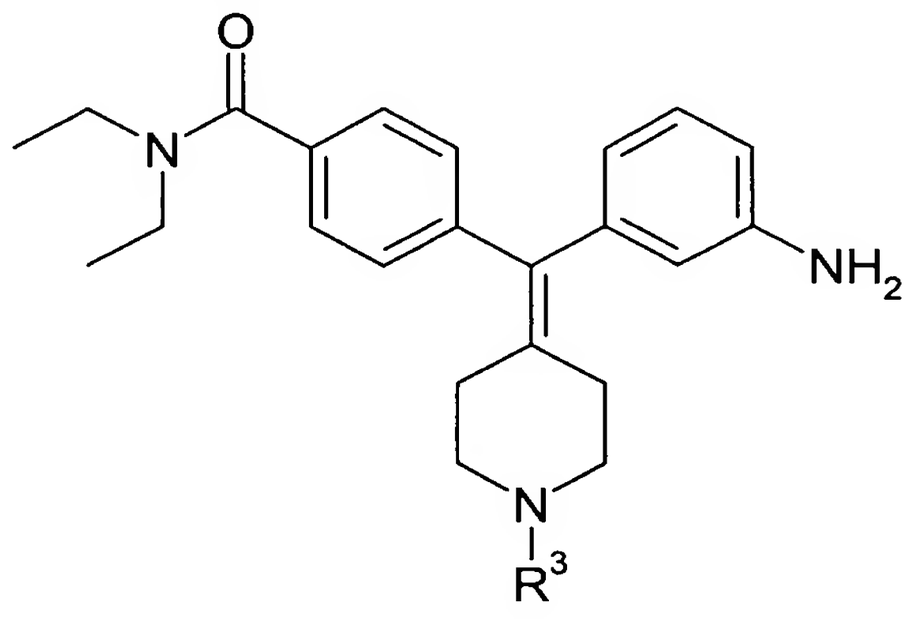
R^3 is selected from C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl, wherein said C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-6}alkyl, halogenated C_{1-6}alkyl, -NO_2, -CF_3, C_{1-6}alkoxy and halogen.

22. (Original) A process for preparing a compound of formula VIII,



VIII

comprising: reacting a compound of formula VII,



VII

with R^8-Z to form the compound of formula VIII:

wherein

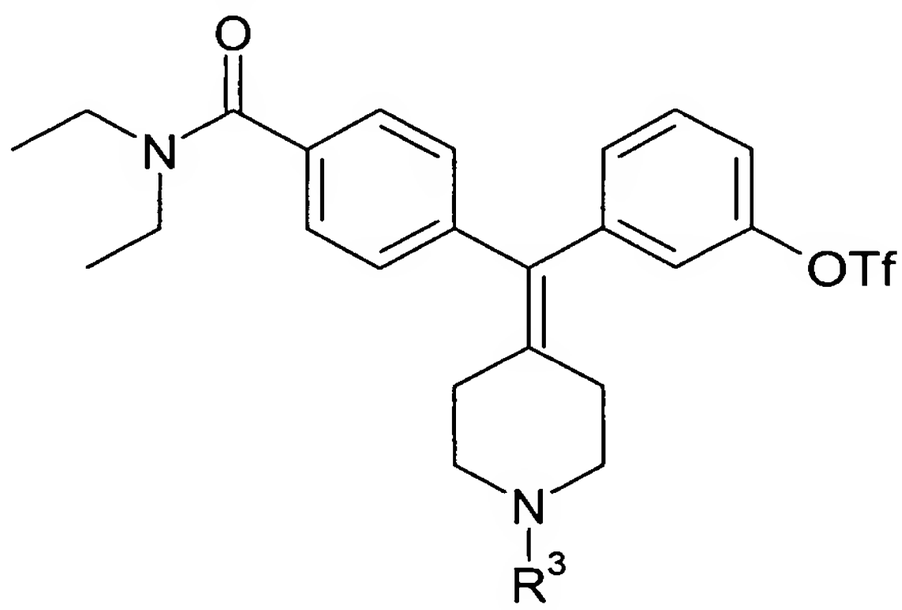
Z is selected from –NCO and -NCS;

Y is selected from –C(=O)NH- and –C(=S)NH-;

R^8 is selected from C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-10}cycloalkyl, and C_{3-10}cycloalkyl-C_{1-4}alkyl; wherein said C_{3-6}alkyl, C_{6-10}aryl, C_{2-6}heteroaryl, C_{6-10}aryl-C_{1-4}alkyl, C_{2-6}heteroaryl-C_{1-4}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with C_{1-4}alkyl, halogen, -CF_3, -OH, C_{1-3}alkoxy, phenoxy, and halogen; and

R^3 is selected from C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl, wherein said C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-6}alkyl, halogenated C_{1-6}alkyl, -NO_2, -CF_3, C_{1-6}alkoxy and halogen.

23. (Original) A compound of formula V,



V

wherein

R^3 is selected from C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl, wherein said C_{1-6}alkyl-O-C(=O)-, C_{1-6}alkyl, C_{3-6}cycloalkyl, and C_{3-6}cycloalkyl-C_{1-4}alkyl are optionally substituted with one or more groups selected from C_{1-6}alkyl, halogenated C_{1-6}alkyl, -NO_2, -CF_3, C_{1-6}alkoxy and halogen.